

PCT

WORLD INTELLECTUAL PROPERTY ORGANIZATION  
International Bureau



INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification 6 :  C08J 7/04		A1	(11) International Publication Number: <b>WO 98/58989</b>  (43) International Publication Date: 30 December 1998 (30.12.98)
(21) International Application Number: PCT/DK98/00264  (22) International Filing Date: 19 June 1998 (19.06.98)  (30) Priority Data: 0730/97 20 June 1997 (20.06.97) DK		(81) Designated States: AL, AM, AT, AT (Utility model), AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, CZ (Utility model), DE, DE (Utility model), DK, DK (Utility model), EE, EE (Utility model), ES, FI, FI (Utility model), GB, GE, GH, GM, GW, HU, ID, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SK (Utility model), SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZW, ARIPO patent (GH, GM, KE, LS, MW, SD, SZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG).  Published <i>With international search report.</i>	
(71) Applicant (for all designated States except US): COLOPLAST A/S [DK/DK]; Holtedam 1, DK-3050 Humlebaek (DK).  (72) Inventor; and  (75) Inventor/Applicant (for US only): MADSEN, Niels, Joergen [DK/DK]; Forelvej 18, DK-3450 Alleroed (DK).  (74) Common Representative: COLOPLAST A/S; Patent Dept., Attn.: Kim Nilausen, Holtedam 1, DK-3050 Humlebaek (DK).			
(54) Title: A HYDROPHILIC COATING AND A METHOD FOR THE PREPARATION THEREOF  (57) Abstract  A hydrophilic coating comprising a polymer containing a hydrophilic prepolymer containing reactive sites cross-linked through polymerisation of vinylic unsaturated groups, said coating comprising a water soluble compound shows a significantly increased water retention and lower friction coefficient when wet.			

**FOR THE PURPOSES OF INFORMATION ONLY**

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AL	Albania	ES	Spain	LS	Lesotho	SI	Slovenia
AM	Armenia	FI	Finland	LT	Lithuania	SK	Slovakia
AT	Austria	FR	France	LU	Luxembourg	SN	Senegal
AU	Australia	GA	Gabon	LV	Latvia	SZ	Swaziland
AZ	Azerbaijan	GB	United Kingdom	MC	Monaco	TD	Chad
BA	Bosnia and Herzegovina	GE	Georgia	MD	Republic of Moldova	TG	Togo
BB	Barbados	GH	Ghana	MG	Madagascar	TJ	Tajikistan
BE	Belgium	GN	Guinea	MK	The former Yugoslav Republic of Macedonia	TM	Turkmenistan
BF	Burkina Faso	GR	Greece			TR	Turkey
BG	Bulgaria	HU	Hungary	ML	Mali	TT	Trinidad and Tobago
BJ	Benin	IE	Ireland	MN	Mongolia	UA	Ukraine
BR	Brazil	IL	Israel	MR	Mauritania	UG	Uganda
BY	Belarus	IS	Iceland	MW	Malawi	US	United States of America
CA	Canada	IT	Italy	MX	Mexico	UZ	Uzbekistan
CF	Central African Republic	JP	Japan	NE	Niger	VN	Viet Nam
CG	Congo	KE	Kenya	NL	Netherlands	YU	Yugoslavia
CH	Switzerland	KG	Kyrgyzstan	NO	Norway	ZW	Zimbabwe
CI	Côte d'Ivoire	KP	Democratic People's Republic of Korea	NZ	New Zealand		
CM	Cameroon			PL	Poland		
CN	China	KR	Republic of Korea	PT	Portugal		
CU	Cuba	KZ	Kazakhstan	RO	Romania		
CZ	Czech Republic	LC	Saint Lucia	RU	Russian Federation		
DE	Germany	LI	Liechtenstein	SD	Sudan		
DK	Denmark	LK	Sri Lanka	SE	Sweden		
EE	Estonia	LR	Liberia	SG	Singapore		

**TITLE**

A hydrophilic coating and a method for the preparation thereof

**FIELD OF THE INVENTION**

The present invention relates to a hydrophilic coating. Furthermore, the invention 5 relates to a medical device provided with such a hydrophilic coating and a method for providing a medical device or other product with a hydrophilic coating as well as the use of a water soluble compound in the preparation of a medical device or instrument comprising a hydrophilic coating being crosslinked.

**BACKGROUND OF THE INVENTION**

10 It is known to coat medical devices, e.g. catheters for introduction into human cavities such as blood vessels, digestive organs and the urinary system, with a hydrophilic coating, normally as a minimum applied on that part of the surface which is introduced or comes into contact with mucous membranes, etc., during introduction of the device. Whereas such coating is not particularly smooth when 15 dry, so that the handling of the device does not become inconvenient, it becomes extremely slippery when it is swelled with water, preferably immediately before introduction into the human body and thus ensures a substantially painless introduction with a minimum of damage on tissue.

A large number of methods are known for the production of hydrophilic surface 20 coatings.

These methods are mainly based on the fact that the substrate to be provided with a hydrophilic surface coating, in the course of one or more process stages with intermediary drying and curing, is coated with one or more (mostly two) layers, which are brought to react with one another in various ways, e.g. by polym- 25 erisation initiated by irradiation, by graft polymerisation, by the formation of interpolymeric network structures, or by direct chemical reaction. Known hydrophilic coatings and processes for the application thereof are e.g. disclosed in Danish Patent No. 159,018, published European Patent Application Nos.

EP 0 389 632, EP 0 379 156, and EP 0 454 293, European Patent No.  
EP 0 093 093 B2, British Patent No. 1,600,963, US Patent Nos. 4,119,094,  
4,373,009, 4,792,914, 5,041,100 and 5,120,816, and into PCT Publication Nos.  
WO 90/05162 and WO 91/19756.

5 According to a method disclosed in US Patent No. 5,001,009, a hydrophilic surface coating is prepared on a substrate by applying, in two stages or in one combined stage, on the substrate a reactive or an adhesive primer layer and then the actual hydrophilic surface layer which, in this case, comprises polyvinylpyrrolidone [PVP] as the active constituent. By this method, no chemical reaction  
10 takes place between the components of the two layers applied.

Where a device of said type, e.g. a catheter for intermittent catherisation, is to remain inside the body only for a short period, there may be a risk that water will be extracted from the hydrophilic surface coating and into the body fluids in the surrounding mucous membranes etc., owing to the higher osmotic potential of said  
15 body fluids. As a result of the extraction of water, the hydrophilic surface coating will have a tendency to become less slippery and to stick to surrounding tissues, and the removal of the medical device from the body may cause pain or damage the tissue. This is especially a problem when carrying out urodynamic examinations via a catheter.

20 European Patent No. EP 0 217 771 describes a method of forming a hydrophilic coating in order to retain the slipperiness for a longer period of time by applying a non-reactive hydrophilic polymer surface layer to a substrate, applying to the non-reactive hydrophilic surface polymer a solution comprising a solvent and above 2% (weight per volume) of an osmolality-increasing compound selected  
25 from the group consisting of mono and disaccharides, sugar alcohols, and non-toxic organic and inorganic salts, with the proviso that the osmolality-increasing compound is not a trihalogenide such as  $KI_3$  ( $KI/I_2$ ), and evaporating the solvent.

International patent publication No. WO 94/16747 discloses a hydrophilic coating with improved retention of water on a surface, especially a surface of a medical device such as a urethra catheter, prepared by applying to the surface in one or more process steps at least one solution of components that will combine to form 5 the hydrophilic coating. During the final step, the surface is coated with an osmolality promoting agent which is dissolved or emulgated in the solution or in the last solution to be applied, forming the hydrophilic coating. WO 94/16747 does not disclose cross-linked coatings.

WO 89/09246 discloses solid shaped structures having a surface coated with 10 crosslinked hydrophilic polymer, the coating being durable and exhibiting a low coefficient of friction when wet. It is stated that the degree of crosslinking is critical and is to be controlled by the operating conditions chosen as too much crosslinking reduces or completely eliminates the low friction surface property, and too little crosslinking negatively affects the durability of the coating. WO 15 89/09246 does not disclose the presence of a water soluble or osmolality-increasing compound in the coating.

Although a hydrophilic coating including an osmolality-increasing agent shows improved properties as compared with conventionally prepared hydrophilic surface coatings, there is still a need of medical devices, especially catheters with a 20 hydrophilic coating conserving the slipperiness for a longer period of time in order to ensure that the coating has not lost its effect when e.g. a catheter is to be retracted.

#### BRIEF DESCRIPTION OF THE INVENTION

It has surprisingly been found that a medical device or instrument having a cross-linked hydrophilic coating comprising a water soluble compound shows a significant increased water retention and a significant decrease of the friction coefficient against living tissue as compared to a device or instrument having a cross-linked hydrophilic coating without a water soluble compound.

Thus, the present invention relates to a hydrophilic coating comprising a covalently cross-linked hydrophilic polymer and a water soluble compound and to a method for the preparation thereof. Furthermore, the invention relates to a medical device provided with such a hydrophilic coating and a method for providing a 5 medical device or other product with a hydrophilic coating comprising a cross-linked hydrophilic polymer and a water soluble compound as well as the use of a water soluble compound in the preparation of a medical device or instrument comprising a cross-linked hydrophilic coating.

#### DETAILED DESCRIPTION OF THE INVENTION

10 In its broadest aspect, the invention relates to a hydrophilic coating comprising a cross-linked hydrophilic polymer and a water soluble agent.

Thus, it has been found that crosslinked hydrophilic coating systems containing a low molecular weight water soluble organic or inorganic compound show a significant increase in water retention and a significant decrease in friction coefficient 15 against living tissue as compared to non-cross-linked or physically cross-linked hydrophilic polymers of the same types containing osmolality increasing ingredients.

Generally, cross-linked coatings show a higher abrasion resistance as compared to the non-cross-linked or physically cross-linked hydrophilic coatings. Without 20 restricting the invention to any specific hypothesis it is assumed that the very low friction coefficients of the coatings of the invention as compared to existing non-cross-linked or physically cross-linked hydrophilic coatings as well as the higher durability of the coatings of the invention is to be ascribed to the fact that the coating of the invention is covalently crosslinked and that the water soluble compound 25 is readily dissolved and leached out of the coating when wetting the same before use leaving interspaces in the crosslinked hydrophilic coating comprising water physically bound in the coating. Thus, the coating shows a very low friction and the water only slowly leaves the coating which contributes to the conservation of the slipperiness of the coating.

It is especially preferred that the cross-linking is effected through polymerisation of vinylic unsaturated groups as such cross-linking is relatively simple to control and has proven to produce hydrophilic coatings having a low friction coefficient and high durability. Other preferred cross-linking may be obtained using polycarboxyl or polyhydroxyl compounds such as polyacrylic acid or polyethylene glycols and optionally isocyanates forming cross-linked polyurethanes

Cross-linking of hydrophilic coatings according to the invention may be effected by radiation and is preferably effected by activation using UV light.

The water soluble compound used according to the invention may be any compound that ensures the desired slipperiness of the hydrophilic coating when moistened.

The water soluble compound used according to the invention may preferably be selected from mono, di and oligosaccharides, sugar alcohols, polypeptides, non toxic organic salts, inorganic salts, organic acids, inorganic acids, urea and mixtures hereof.

Thus, the water soluble compound may be selected from glucose, sorbitol, halides, nitrates, acetates, citrates or benzoates of alkali metals or alkaline earth metals or silver such as sodium chloride, sodium citrate, sodium benzoate, calcium chloride, potassium chloride, potassium iodide or potassium nitrate; acetic acid, amino acids such as glycine and urea. The compound or mixture of compound is preferably non-toxic and may be soluble or insoluble in the solution into which the compound is incorporated. It is preferred when the osmolality-increasing compound or mixture may be incorporated into a solution by dissolution or emulsification.

Especially preferred are hydrophilic coatings according to the invention comprising sodium chloride or urea.

In hydrophilic coatings according to the invention sodium chloride or urea may be used in an amount of 0.1 - 200 % (w/w) on the basis of the dry hydrophilic coating giving rise to hydrophilic coatings showing even lower friction coefficient than that of known hydrophilic coatings comprising an "osmolality-increasing compound". Preferably, sodium chloride or urea is present in an amount of 1 - 80 % (w/w), more preferred in an amount from 2 to 30 % (w/w).

The cross-linked hydrophilic coating may be any cross-linked coating.

The cross-linked hydrophilic coating may e.g. be of the type disclosed in European patent application No. EP 0 289 996 A2. Such a coating is formed and applied to a medical device or instrument in the form of a solution containing a water-soluble polymer, more particularly polyvinylpyrrolidone or a copolymer thereof, one or more radically polymerisable vinyl monomers and a photo initiator and the applied solution is exposed to an UV radiation for curing purposes. The osmolality-increasing compound is incorporated in the solution applied to the medical device or instrument.

The cross-linked hydrophilic coating may also comprise a cross-linked hydrophilic polymer wherein the polymer comprises a prepolymer containing reactive sites cross-linkable through polymerisation of vinylic unsaturated groups and optionally one or more saturated polymers crosslinked through vinylic groups. In this case, the water soluble compound is also incorporated in a sol or solution of a hydrophilic prepolymer containing reactive sites cross-linkable through polymerisation of vinylic unsaturated groups and optionally one or more hydrophilic polymers to be coated onto the medical device and cross-linked by activation through UV-light or radiation and optionally hydrolysed and optionally neutralised.

The cross-linked hydrophilic coating may also be a hydrophilic coating comprising a cross-linked polyvinylpyrrolidone or a copolymer of N-vinylpyrrolidone and optionally a photo initiator.

A hydrophilic coating according to the invention may be used for coating the surface or a part thereof of a wide range of products in order to impart give the surface a low friction. As examples of products which may be provided with a surface having a low friction when wet are medical instruments such as catheters, endo and laryngoscopes, tubes for feeding, or drainage or endotracheal use, condoms, barrier coatings, e.g. for gloves, wound dressings, contact lenses, implantates, extracorporeal blood conduits, membranes e.g. for dialysis, blood filters, devices for circulatory assistance, packaging for foodstuff, razor blades, fishermen's net, conduits for wiring, water pipes having a coating inside, sports articles, cosmetic additives, mould release agents, and fishing lines and nets.

Furthermore, the invention relates to a medical device or instrument provided with a hydrophilic coating comprising a cross-linked hydrophilic polymer, said coating comprising a water-soluble compound.

The invention especially relates to catheters for intermittent catherisation provided with such a cross-linked hydrophilic coating.

The medical device of the invention is preferably a catheter. The components of a cross-linked hydrophilic coating may be applied onto a catheter made from PVC or polyurethane.

In accordance preferred embodiment of the invention the coatings comprise an antibacterial agent such as a silver salt, e.g. silver sulphadiazine, an acceptable iodine source such as povidone iodine (also called polyvinylpyrrolidone iodine), chlorhexidine salts such as the gluconate, acetate, hydrochloride or the like salts or quaternary antibacterial agents such as benzalkonium chloride or other anti-septics or antibiotics. Antibacterial agents reduces the risk of infection, especially when performing urodynamic examinations.

Still further, the invention relates to a method of producing a medical device or instrument having a hydrophilic coating of a cross-linked hydrophilic polymer,

said coating comprising a water soluble compound, in which method is applied, in one or more steps, a solution of a hydrophilic polymer or prepolymer and optionally a monomer, oligomer or polymer and a water soluble compound, the solvent is evaporated and the coating is cross-linked by activation through radiation 5 and optionally hydrolysing and optionally neutralising the hydrophilic coating.

The method according to the invention may be carried out in a manner where a first primer layer is first applied to the substrate. This primer layer may for instance be a cross-linked primer layer, e.g. based on PVP, or a non-cross-linked primer layer based on nitro-cellulose applied in a solution. After drying of the 10 primer layer, an outer layer is applied consisting of a solution of polyvinylpyrrolidone in a solvent selected from among tetrahydrofuran, methyl(ene) chloride, toluene, acetone, a lower aliphatic alcohol, cyclohexanone, C<sub>2</sub>-C<sub>4</sub>-alkyl acetates, butyrolactone, and dimethylformamide, the most important constituent being ethyl alcohol. According to the invention, this solution contains the water soluble 15 compound.

The term "urea" used herein should be understood to comprise urea as well as which has been N-substituted or N,N-disubstituted by lower alkyl.

"Lower alkyl" is used in the present context to designate straight or branched or cyclic aliphatic groups having from 1 to 6 carbon atoms, preferably from 1 to 4 20 carbon atoms such as methyl, ethyl, propyl, isopropyl or n, iso or tert.butyl.

## MATERIALS AND METHODS

### **Method for Determination of the Friction and Water Retension.**

The Standard Test Method for Static and Kinetic Coefficient of Friction of Plastic Film and Sheeting, ASTM D 1894 - 93 was modified for testing the friction coefficient and wear on plastic tubes and catheters.

The tubes or catheters were cut in lengths of 10 cm and fixed on a stainless steel plate with two stainless steel rods as shown in ASTM D 1894 - 93. The rods had diameters comparable with the inner diameter of the tubes or catheters to keep their shape even when heavy sledges were placed upon them.

5

The friction was determined under water or after wetting by dipping in water for 1 minute. The pulling force from the sledge was measured in Newtons.

Water retension of the coatings was determined by measuring the friction for up to 16 or 18 minutes after dipping the tubes in water for determining the time to 10 reach dryness.

Polyvinylpyrrolidone: Plasdone® K90 Povidone USP from International Speciality Products.

Ethanol: Absolute Alcohol.

Butyrolactone: Gamma-butyrolactone from International Speciality Products.

15 UV catalyst: ESACURE KIP 150 from Lamberti SpA.

The invention is further explained in detail with reference to the below working examples disclosing embodiments of the invention. The embodiments are illustrative of the principles of the invention and it is clear to the skilled in the art that modifications may be made without deviating from the gist of the invention, the 20 scope of which is set forth in the appended claims.

## EXPERIMENTAL PART

### EXAMPLE A

Preparation of a catheter having a cross-linked hydrophilic PVP coating.

4.9 parts of PVP K 90 and 0.1 parts ESACURE KIP 150 was dissolved in 95  
5 parts of an ethanol/gamma butyrolactone (85/15) solvent mixture and used as a  
primer coat. PVC-catheters were dipped in the primer solution and dried in an  
oven for 1 minute at 60°C before dipping in a solution of 4 parts of PVP K 90 dis-  
solved in 96 parts of an ethanol/gamma butyrolactone (85/15) solvent mixture.  
After drying for 30 minutes the PVC-catheters were exposed to UV-light with a  
10 wave length between 200 and 300 nm for 5 minutes.

### EXAMPLE B

Preparation of a catheter having a hydrophilic coating.

PVC-catheters were dipped in a primer solution of 4 parts of a medical grade  
thermoplastic polyurethane and 2 parts of nitrocellulose dissolved in 94 parts of  
15 THF and afterwards dried in an oven for 15 minutes at 60°C.  
4.0 parts of PVP K 90 was dissolved in 96 parts of an ethanol/gamma butyrolac-  
tone (85/15) solvent mixture and coated onto the PVC-catheters and dried 1 hour  
in an oven at 60°C.

### EXAMPLE 1.

20 Preparation of a catheter according to the invention having a cross-linked hydro-  
philic coating comprising urea.

PVC-catheters were dipped in the primer solution as made in Example A, after  
drying 1 minute at 60°C the catheters were dipped in a topcoat solution of 3.36  
parts of PVP K 90 and 0.64 parts of urea dissolved in 96 parts of an  
25 ethanol/gamma butyrolactone (85/15) solvent mixture. The coating was further

dried in an oven for 30 minutes at 60°C and exposed to UV-light with a wave length between 200 and 300 nm for 5 minutes.

#### **COMPARATIVE EXAMPLE 1**

Preparation of a catheter having a non-crosslinked hydrophilic coating comprising urea.

PVC-catheters were dipped in the PU/nitrocellulose primer solution as made in comparative Example B and dried for 15 minutes before they were dipped in the PVP-solution containing 3.36 parts of PVP K 90, 0.64 parts of urea and 96 parts of an ethanol/gamma butyrolactone (85/15) solvent mixture. The catheters were further dried 1 hour.

#### **EXAMPLE 2**

Preparation of a catheter according to the invention having a cross-linked hydrophilic coating comprising sodium chloride.

PVC-catheters were dipped in the primer solution as made in Example A. and 15 dried for 1 minute at 60°C. Then the catheters were dipped in the PVP-solution of 3.6 parts of PVP K 90 and 0.4 parts sodium chloride dissolved in 96 parts of an ethanol/gamma butyrolactone/water (64/20/16) solvent mixture. The catheters were further dried for 30 minutes and exposed to UV-light with a wave length range between 200 and 300 nm. for 5 minutes.

#### **20 COMPARATIVE EXAMPLE 2**

Preparation of a catheter having a non-crosslinked hydrophilic coating comprising sodium chloride.

PVC-catheters were dipped in the PU/nitrocellulose primer solution as made in the comparative Example B and dried for 5 minutes before they were dipped in 25 the PVP-solution containing 3.6 parts of PVP K 90 and 0.4 parts sodium chloride

dissolved in 96 parts of an ethanol/gamma butyrolactone/water (64/20/16) solvent mixture.

The result of determination of the friction force of the 6 hydrophilic coatings is shown in table 1. The figures are means of 3 readings.

5 Table 1

Friction force measured on cross-linked and non cross-linked PVP coatings and cross-linked and non cross-linked PVP coatings comprising low molecular weight compounds, urea or sodium chloride.

Example	A	B	1	Comp 1	2	Comp 2
Initial reading (N)	0.10	0.07	0.03	0.07	0.10	0.3
After 2 min (N)	0.11	0.08	0.03	0.07	0.10	0.7
After 4 min (N)	dry	0.22	0.03	0.08	0.11	dry
After 6 min (N)		dry	0.03	0.10	0.3	
After 8 min (N)			0.04	dry	dry	
After 10 min (N)			0.04			
After 12-18 min (N)			dry			

It appears from the table that the coatings of the invention are superior with respect to retaining low friction as compared to coatings not being cross-linked and coatings not being cross-linked but comprise an osmolality increasing agent.

### EXAMPLE 3

Preparation of a tube having a hydrophilic coating according to the invention.

A dispersion of polyisocyanate was prepared from 50 parts of a 15 PVP/hydroxyethylmethacrylate-copolymer having 10 mole % HEMA, 10 parts polyethyleneglycol having  $M_w=10,000$ , 30 parts of polyisocyanate (Bayhudur® VP LS 2032 from Bayer AG) and 10 parts of sodium chloride in the form of a 10% dispersion in water. Tubes of polyurethane were dipped into the dispersion and dried at 70°C for three hours in an oven. The coating showed a low friction coefficient and high abrasion resistance.

**EXAMPLE 4**

Leaching of water soluble compound from coating of the invention.

A catheter having a coating of 5 microns dry thickness according to the invention prepared according to Example 1 was tested for leaching of urea in water after 5 30 seconds and 24 hours.

The catheter was dipped in 60 milliliters of water and the concentration of urea was determined by a spectrophotometric method after 30 seconds and 24 hours respectively. The coating swells to about 80 microns. It is assumed that all urea had been leashed from the coating after 24 hours. The concentration of urea in 10 the liquid was 0.0348 grams per liter after 30 seconds and 0.0373 grams per liter after 24 hours. Thus, 93.3% of the urea is leached out of the coating after 30 seconds and the urea remaining in the coating corresponds to 0.06% of the inclusion water of the coating.

Based on the results it is demonstrated that the coating of the invention has a 15 low friction coefficient although being hypotonic.

**CLAIMS**

1. A hydrophilic coating comprising a covalently cross-linked hydrophilic polymer, said coating comprising a water soluble compound.
2. A hydrophilic coating as claimed in claim 1, characterised in that the hydrophilic polymer is cross-linked via vinylic groups.
3. A hydrophilic coating as claimed in claim 1 or 2, characterised in that the water soluble compound is selected from mono, di and oligosaccharides, sugar alcohols, polypeptides, non toxic organic salts, inorganic salts, organic acids, inorganic acids, urea and mixtures hereof.
- 10 4. A hydrophilic coating as claimed in claim 3, characterised in that the water soluble compound is selected from glucose, sorbitol, halides, nitrates, acetates, citrates or benzoates of alkali metals or alkaline earth metals or silver such as sodium chloride, sodium citrate, sodium benzoate, calcium chloride, potassium chloride, potassium iodide or potassium nitrate; acetic acid, glycine and urea.
- 15 5. A hydrophilic coating as claimed in any of claims 1 - 4, characterised in that the water soluble compound is present in an amount of 0.1 - 200 % (w/w) on the basis of the hydrophilic coating.
6. A hydrophilic coating as claimed in claim 5, characterised in that the water soluble compound is present in an amount of 2 - 80 % (w/w).
- 20 7. A hydrophilic coating as claimed in any of claims 1 - 6, characterised in that the water soluble compound is urea.
8. A hydrophilic coating as claimed in any of claims 1-7, characterised in that the cross-linked coating contains an antibacterial agent.

9. A medical device or instrument having a hydrophilic coating comprising a cross-linked hydrophilic polymer, said coating comprising a water soluble compound.
10. A method of producing a medical device or instrument having a hydrophilic coating comprising a cross-linked hydrophilic polymer, said coating comprising a water soluble compound, characterised in applying, in one or more steps, a solution of a hydrophilic prepolymer and optionally a monomer, oligomer or polymer and a water soluble compound, evaporating the solvent and cross-linking the coating by activation through radiation and optionally hydrolysing and optionally neutralising the hydrophilic coating.

## INTERNATIONAL SEARCH REPORT

International application No.  
PCT/DK 98/00264

## A. CLASSIFICATION OF SUBJECT MATTER

IPC6: C08J 7/04

According to International Patent Classification (IPC) or to both national classification and IPC

## B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC6: C08J, A61L

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

SE, DK, FI, NO classes as above

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

WPI, REGISTRY, CAPLUS

## C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	WO 9416747 A1 (COLOPLAST A/S), 4 August 1994 (04.08.94), abstract, claims --	1-10
Y	WO 8909246 A1 (E.I. DU PONT DE NEMOURS AND COMPANY), 5 October 1989 (05.10.89), abstract, page 4, line 18 - line 34; page 5, line 11 - line 15; page 6, line 25 - page 7, line 12; page 11, line 34 - page 12, line 15; exemple 4; claims 1-2, 21-24, 36, 41 --	1-10
A	EP 0289996 A2 (WILKINGSON SWORD GESELLSCHAFT MIT BESCHRÄNKTER HAFTUNG), 9 November 1988 (09.11.88), column 3, line 23 - line 50 -----	1-2

 Further documents are listed in the continuation of Box C. See patent family annex.

- \* Special categories of cited documents:
- "A" document defining the general state of the art which is not considered to be of particular relevance
- "B" earlier document but published on or after the international filing date
- "C" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- "D" document referring to an oral disclosure, use, exhibition or other means
- "P" document published prior to the international filing date but later than the priority date claimed
- "T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
- "X" document of particular relevance: the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
- "Y" document of particular relevance: the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
- "&" document member of the same patent family

Date of the actual completion of the international search

28 August 1998

Date of mailing of the international search report

14-09- 1998

Name and mailing address of the ISA

Swedish Patent Office  
Box 5055, S-102 42 STOCKHOLM  
Facsimile No. + 46 8 666 02 86

Authorized officer

Sofia Nikolopoulou  
Telephone No. + 46 8 782 25 00

**INTERNATIONAL SEARCH REPORT**

Information on patent family members

International application No.

27/07/98

PCT/DK 98/00264

Patent document cited in search report	Publication date	Patent family member(s)		Publication date
WO 9416747 A1	04/08/94	AU	660873 B	06/07/95
		AU	5880694 A	15/08/94
		DK	7193 A,B	22/07/94
		EP	0639990 A	01/03/95
<hr/>				
WO 8909246 A1	05/10/89	AU	629203 B	01/10/92
		AU	3219889 A	16/10/89
		DK	228690 A	21/09/90
		EP	0418247 A	27/03/91
		JP	3503379 T	01/08/91
<hr/>				
EP 0289996 A2	09/11/88	AU	604883 B	03/01/91
		AU	1564688 A	10/11/88
		CA	1328989 A	03/05/94
		DE	3814135 A	24/11/88
		DE	3866008 A	12/12/91
		HK	62795 A	05/05/95
		IN	172419 A	17/07/93
		JP	2631997 B	16/07/97
		JP	63294971 A	01/12/88
		US	5005287 A	09/04/91